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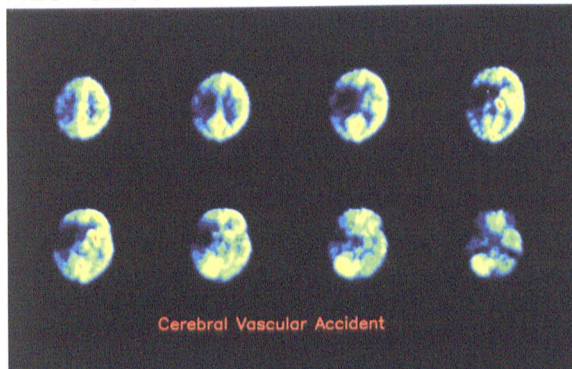
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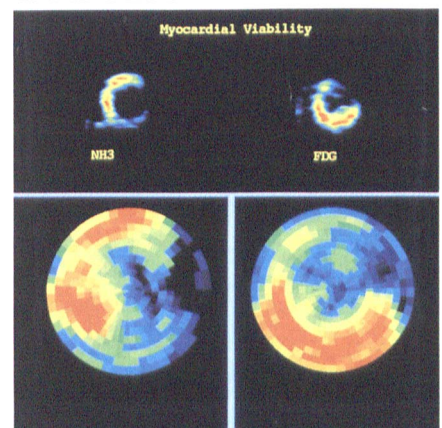
NEUROLOGY



"PET has the ability to measure biochemical responses to disease in the brain prior to gross changes in anatomy and, in some cases, prior to symptom onset resulting in early diagnosis and improved patient management."

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Vice-Chairman of Neurology
Professor of Neurology and Radiology
UCLA School of Medicine

CARDIOLOGY

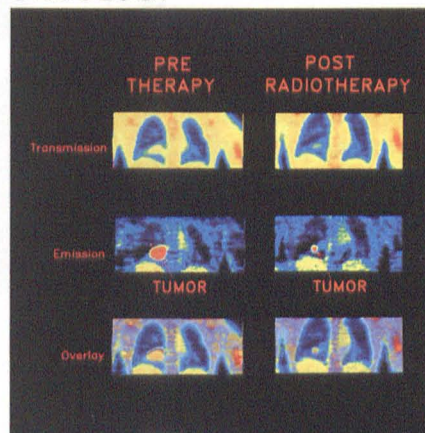


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"PET provides unique non-invasive information on behavior, treatment response, and recurrence rate of solid tumors. Clinical PET promises to greatly impact the practice of oncology."

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EDITORIAL COMMUNICATIONS should be sent to the Editor: H. William Strauss, MD, The Journal of Nuclear Medicine, Room 5406 MGH-East, Bldg. 149, 13th St., Charlestown, MA 02129 (617) 726-5786. *Books and monographs* covering the use of nuclear medicine and its allied disciplines will be reviewed as space is available. *Send review copies to the Editor.*

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SUBSCRIPTION RATES for 1992 calendar year are \$120 within the United States; \$130 for Canada and Pan American countries; \$160 elsewhere. Student subscriptions are \$70 (with proof of student status). Single copies \$10.00; foreign \$11.00; convention issue (May) \$12.00; foreign \$13.00. Make checks payable, in U.S. dollars drawn on U.S. banks, to The Society of Nuclear Medicine. Notify the Society of change of address and telephone number at least 30 days before date of issue by sending both the old and new addresses.

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Reality

Cogito, ergo sum (I think, therefore I am).
Rene Descartes

A realist lets circumstances decide which end of the telescope to look through.
Anon

Vision is the art of seeing things invisible.
Jonathan Swift

Perception and reality are not two peas in a pod.

Reality *is*.

The reality we perceive not only *is*, but is dependent on circumstances which vary from moment to moment. Given our limited sensorium, what simply *is*, is quickly transmogrified by a multi-tasking mind. To translate the what 'is' into something we can understand, the sensory input is projected on the canvas of the emotional background of the moment. Or at least, that's how I perceive it.

The system is convoluted. Yet the system has an evolutionary advantage. If we faced reality head on and simply experienced things as they are, without processing the material, our ability to interpret and analyze would be lost. We would simply be machines, tallying data, devoid of a sense of wonder, and disinterested in comparative concepts such as beauty and joy.

Of course, looking at this from another angle, our inability to refrain from colorizing reality makes it possible to ask about the actual basis of our own reality. As Alan Watts suggests, do we only see white because there is black, or good because there is bad? To discern the truth, we need to collect many samples and interview many witnesses. After careful analysis, we may approach truth asymptotically.

An awareness of our subjective state is not enough, just as one carefully constructed experiment is never enough to reveal the complete picture. As in the fable of the blind man and the elephant, many results must be knit together to first begin to understand what we are actually seeing, or feeling.

So it is in science, and so it is in publishing.

To help us understand that the reality of Nuclear Medicine, as you see it, is similar to the way we perceive it, we seek your help. In this month's issue of the *Journal*, there is a reader survey form. If you could take a few moments to fill it out, and send or FAX your perceptions to us, we will continue to try and make the *Journal's* realities more closely resemble your expectations.

Of course expectations are the enemy of perception. But that's another story...

H. William Strauss
Editor, *The Journal of Nuclear Medicine*

NRC REQUIREMENT:

"A licensee shall survey for removable contamination, once each week, all areas where radiopharmaceuticals are routinely prepared for use, administered or stored."

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Cyclic Oral Phosphate and Editronate Increase Femoral and Lumbar Bone Mineral Density and Reduce Lumbar Spine Fracture Rate Over Three Years

Forty post-menopausal women with osteoporosis were treated with a sequential, cyclic regimen of oral phosphate for 3 days, etidronate for 2 wk, and calcium salt for 13 wk. The cycle was repeated for 3 yr and the patients were rescanned after every other cycle. Page 1

Myocardial Emission Computed Tomography with Iodine-123-Labeled Beta-Methyl Branched Fatty Acid in Patients with Hypertrophic Cardiomyopathy

Iodine-123-labeled BMIPP SPECT studies were compared to ^{201}Tl in 17 patients with hypertrophic cardiomyopathy. Page 6

Safety and Role of Repeated Administration of Indium-111-Labeled Anti-carcinoembryonic Antigen Monoclonal Antibody ZCE 025 in the Postoperative Follow-up of Colorectal Carcinoma Patients

Twenty-five patients who had undergone resection for colorectal carcinoma were prospectively studied to evaluate the safety of repeated intravenous administration of labeled murine monoclonal antibody, ZCE 025, and to determine whether tumor recurrence and/distant metastases could be detected by radioimmunosintigraphy when anti-mouse antibodies were present in patients' sera. Page 14

Phase I Trial of Iodine-131-Chimeric B72.3 (Human IgG4) in Metastatic Colorectal Cancer

In a Phase I trial of ^{131}I -labeled chimeric B72.3, 12 patients with metastatic colorectal cancer received 18 mCi/m², 27 mCi/m² and 36 mCi/m² of labeled antibody. Page 23

Editorial: Radiolabeled Monoclonal Antibodies for Cancer Therapy and Diagnosis: Is It Really A Chimera?

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Radiation Dosimetry for Technetium-99m-MAG3, Technetium-99m-DTPA, and Iodine-131-OIH Based on Human Biodistribution Studies

Biokinetics and radiation dose estimates for the renal agents $^{99\text{m}}\text{Tc}$ -MAG3, and ^{131}I -OIH were obtained from studies of healthy human volunteers. Page 33

A Dual-Radioisotope Technique for the Evaluation of Penile Blood Flow During Tumescence

The authors describe a technique for arterial and venous penile blood flow during tumescence using ^{133}Xe and $^{99\text{m}}\text{Tc}$ Page 41

Editorial: Vascular Testing for Impotence.

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Radionuclide Assessment of Penile Corporal Venous Leak Using Technetium-99m-Labeled Red Blood Cells

To evaluate penile corporal venous outflow, a method that utilizes intracorporal injection of Tc-RBC was developed and used in 20 patients with erectile dysfunction. Page 49

Technetium-99m-HMPAO SPECT in the Evaluation of Patients with a Remote History of Traumatic Brain Injury: A Comparison with X-ray Computed Tomography

Fifty-three patients with a remote history of traumatic brain injury were studied by SPECT using $^{99\text{m}}\text{Tc}$ -HMPAO and x-ray computed tomography. Page 52

Diagnosis of Sternal Wound Infection by Technetium-99m-Leukocyte Imaging

The $^{99\text{m}}\text{Tc}$ -leukocyte scans of 29 patients, originally referred to rule out

sternal infections, were retrospectively reviewed to evaluate the efficacy of leukocyte imaging. Page 59

Editorial: Imaging Inflammation: Current Role of Labeled Autologous Leukocytes.

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Left Ventricular Diastolic Function in Systemic Sclerosis: Assessment by Radionuclide Angiography

Twenty-four women with systemic sclerosis and 14 age- and sex-matched controls were studied with radionuclide angiography to assess left ventricular function. Page 68

Influence of Ureteral Status on Kidney Washout During Technetium-99m-DTPA Diuresis Renography in Children

Ureteral images from 42 children referred for suspicion of hydronephrosis were reviewed to assess the influence of the ureter on renal washout during $^{99\text{m}}\text{Tc}$ -DTPA diuresis renography. Page 73

Editorial: Effects of Ureteral Function on Assessment of Hydronephrosis.

. Page 78

Technetium-99m-DTPA Aerosol and Gallium-67 Scanning in Pulmonary Complications of Human Immunodeficiency Virus Infection

Gallium-67 chest scans and $^{99\text{m}}\text{Tc}$ -DTPA aerosol clearance measurements were retrospectively compared with results from fiberoptic bronchoscopy exams of 88 patients infected with HIV. Page 81

Absorbed Radiation Dose to Humans from Technetium-99m-Teboroxime

Radiation dose to humans after intravenous administration of $^{99\text{m}}\text{Tc}$ -teboroxime was derived from tissue distribution data obtained from nine normal volunteers. Organ uptake as a percent of injected dose was measured using quantitative SPECT. Page 88

Myocardial Extraction of Teboroxime: Effects of Teboroxime Interaction with Blood

An isolated perfused rat heart preparation was used to determine whether the interaction of blood with either ^{99m}Tc -teboroxime, ^{99m}Tc -sestamibi, or ^{201}Tl affects the extraction of these myocardial perfusion agents. Page 94

Uptake Kinetics of Technetium-99m-Methoxyisobutylisonitrile and Thallium-201 in Adult Rat Heart Endothelial and Fibroblast-like Cells in Comparison to Myocytes

The net uptake of ^{201}Tl and ^{99m}Tc -sestamibi by cultured rat heart endothelial and fibroblast-like cells and quiescent myocytes was examined to determine their role in the uptake of tracers used for myocardial perfusion imaging Page 102

Uptake Mechanism of Technetium-99m-d,l-HMPAO in Cell Cultures of the Dissociated Postnatal Rat Cerebellum

The accumulation and retention mechanisms of ^{99m}Tc -d,l-HMPAO were investigated in cultures of the dissociated rat cerebellum. Page 108

Unique Scintigraphic Findings of Bile Extravasation in the Presence of Ascites: A Complication of Hepatic Transplantation

A 4-mo-old female liver transplant patient was imaged with ^{99m}Tc -HIDA 6 days after transplantation. Page 115

Reversible Increased Technetium-99m-HMPAO Cerebral Cortical Activity: A Scintigraphic Reflection of Luxuriant Hyperperfusion

This study of a hemiparetic and aphasic patient suggests that evanescent

peripheral cerebral hyperemia may represent beneficial cortical collateralization of the peri-infarct area of a deeper lacunar CVA. Page 117

Clinicopathologic Conferences: A Thallium Scan Goes to Court
. Page 120

Comparison of Left Anterior Oblique, Anterior and Geometric Mean Methods for Determining Gastric Emptying Times

To determine if there were significant differences in gastric emptying time measurements, the authors imaged patients in the anterior, posterior and left anterior oblique views. Linear regressions were then obtained using the anterior, left anterior oblique, and geometric mean data. Page 127

Simultaneous Dual-Isotope SPECT Imaging for the Detection and Characterization of Parathyroid Pathology

A patient study illustrating a simultaneous dual-isotope SPECT imaging method is described. Page 131

Kinetic Behavior of Technetium-99m-HMPAO in the Human Brain and Quantification of Cerebral Blood Flow Using Dynamic SPECT

Using dynamic SPECT and a four-compartment model with five parameters, the kinetic behavior of ^{99m}Tc -HMPAO in the brain was investigated in 11 patients with a variety of brain diseases. Page 135

An Instant Kit Method for Labeling Antimyosin Fab¹ with Technetium-99m: Evaluation in an Experimental Myocardial Infarct Model

Antimyosin Fab¹ was labeled with ^{99m}Tc using this kit method. The result was

compared in murine biodistribution studies and in canine experimental infarct model to ^{111}In -antimyosin Fab-DTPA. Page 144

Imaging of the Human Torso Using Cone-Beam Transmission Computed Tomography Implemented on a Rotating Gamma Camera

The feasibility of high quality, cone-beam transmission CT generated on a rotating gamma camera was investigated in three human subjects and compared to conventional CT. Possible imaging protocols are discussed. Page 150

A Method for Comparing Different Procedures of Estimating Regional Glucose Metabolism Using Fluorine-18-Fluorodeoxyglucose

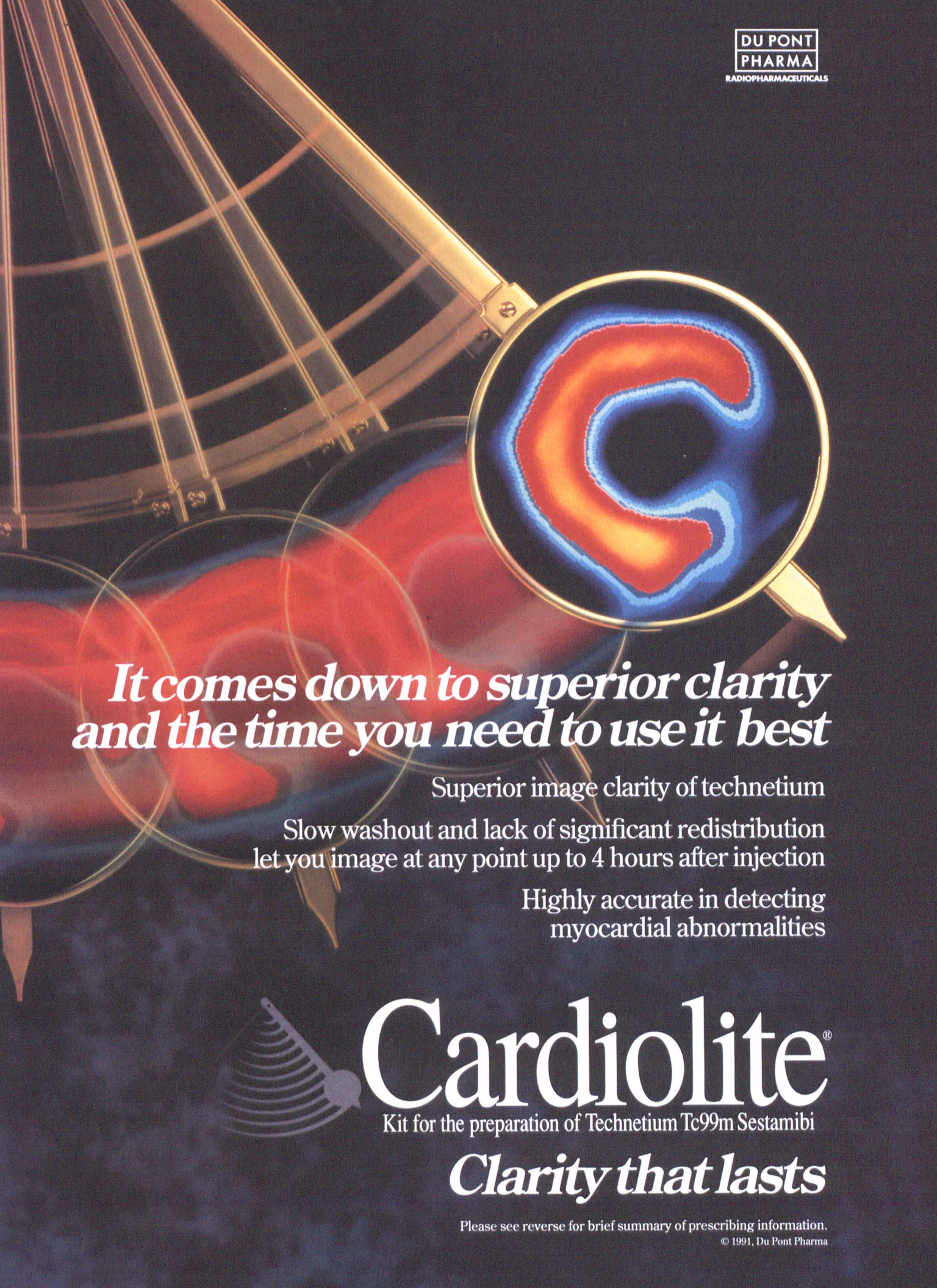
The authors have developed a method based on a simple model of regional cerebral glucose metabolism, allowing for three potential sources of metabolic variability: individual differences in cerebral metabolic rate, consistent regional differences, and error. Page 157

Attenuation Correction for Bremsstrahlung Imaging Using the Gamma Camera

Quantitative imaging of bremsstrahlung from pure beta emitters is proposed as a means for in-vivo management of antibody therapy. Page 161

Commentary: Radiation Safety for Beginners. Page 167

Commentary: The Utility of Single-Photon Absorptiometry and Dual-Energy X-ray Absorptiometry
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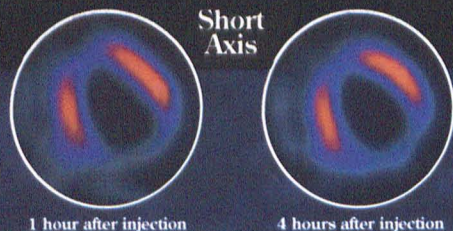
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Please see reverse for brief summary of prescribing information.

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CARDIOLITE scans (SPECT) from a 62-year-old male with three prior myocardial infarctions (LFOV camera equipped with a high-resolution collimator, 64 x 64 matrix, 180° arc RAO to LPO, 64 projections, 25 s/projection).



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Brief Summary

Cardiolite® Kit for the preparation of Technetium Tc99m Sestamibi

F O R D I A G N O S T I C U S E

DESCRIPTION: Each 5 mL vial contains a sterile, non-pyrogenic, lyophilized mixture of:

Tetrakis (2-methoxy isobutyl isonitrile) Copper (I) tetrafluoroborate - 1.0 mg
Sodium Citrate Dihydrate - 2.6 mg
L-Cysteine Hydrochloride Monohydrate - 1.0 mg
Mannitol - 20 mg
Stannous Chloride, Dihydrate, minimum ($\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$) - 0.025 mg
Stannous Chloride, Dihydrate, ($\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$) - 0.075 mg
Tin Chloride (Stannous and Stannic) Dihydrate, maximum (as $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$) - 0.086 mg

Prior to lyophilization the pH is 5.3 to 5.9. The contents of the vial are lyophilized and stored under nitrogen.

This drug is administered by intravenous injection for diagnostic use after reconstitution with sterile, non-pyrogenic, oxidant-free Sodium Pertechnate Tc99m Injection. The pH of the reconstituted product is 5.5 (5.0-6.0). No bacteriostatic preservative is present.

The precise structure of the technetium complex is $\text{Tc99m}[\text{MIBI}]_n^+$ where MIBI is 2-methoxy isobutyl isonitrile.

INDICATIONS AND USAGE: CARDIOLITE®. Kit for the preparation of Technetium Tc99m Sestamibi, is a myocardial perfusion agent that is useful in distinguishing normal from abnormal myocardium, and in the localization of the abnormality, in patients with suspected myocardial infarction. It is also useful in the evaluation of myocardial function using the first-pass technique.

CONTRAINDICATIONS: None known.

WARNINGS: In studying patients in whom cardiac disease is known or suspected, take care to assure continuous monitoring and treatment in accordance with safe, accepted clinical procedure.

PRECAUTIONS:

GENERAL

The contents of the vial are intended only for use in the preparation of Technetium Tc99m Sestamibi and are not to be administered directly to the patient without first undergoing the preparative procedure (as outlined in the full prescribing information).

Radioactive drugs must be handled with care and appropriate safety measures should be used to minimize radiation exposure to clinical personnel. Also, care should be taken to minimize radiation exposure to the patients consistent with proper patient management.

Contents of the kit before preparation are not radioactive. However, after the Sodium Pertechnate Tc99m Injection is added, adequate shielding of the final preparation must be maintained.

The components of the kit are sterile and non-pyrogenic. It is essential to follow directions carefully and to adhere to strict aseptic procedures during preparation.

Technetium Tc99m labeling reactions involved depend on maintaining the stannous ion in the reduced state. Hence, Sodium Pertechnate Tc99m Injection containing oxidants should not be used.

Technetium Tc99m Sestamibi should not be used more than six hours after preparation.

Radiopharmaceuticals should be used only by physicians who are qualified by training and experience in the safe use and handling of radionuclides and whose experience and training have been approved by the appropriate government agency authorized to license the use of radionuclides.

Carcinogenesis, Mutagenesis, Impairment of Fertility

In comparison with most other diagnostic technetium-labeled radiopharmaceuticals, the radiation dose to the ovaries (1.5 rads/30 mCi) is high. Minimal exposure (ALARA) is necessary in women of childbearing capability. (See Dosimetry subsection in DOSAGE AND ADMINISTRATION section.)

The active intermediate, $\text{Cu}(\text{MIBI})_2\text{BF}_4$, was evaluated for genotoxic potential in a battery of five tests. No genotoxic activity was observed in the Ames, CHO/HPRT and sister chromatid exchange tests (all *in vitro*). At cytotoxic concentrations ($\geq 20 \mu\text{g/mL}$), an increase in cells with chromosome aberrations was observed in the *in vitro* human lymphocyte assay. $\text{Cu}(\text{MIBI})_2\text{BF}_4$ did not show genotoxic effects in the *in vivo* mouse micronucleus test at a dose which caused systemic and bone marrow toxicity (9 mg/kg, $>600 \times$ maximal human dose).

Pregnancy Category C

Animal reproduction and teratogenicity studies have not been conducted with Technetium Tc99m Sestamibi. It is also not known whether Technetium Tc99m Sestamibi can cause fetal harm when administered to a pregnant woman or can affect reproductive capacity. There have been no studies in pregnant women. Technetium Tc99m Sestamibi should be given to a pregnant woman only if clearly needed.

Ideally, examinations using radiopharmaceuticals, especially those elective in nature, of a woman of childbearing capability, should be performed during the first few (approximately 10) days following the onset of menses.

Nursing Mothers

Technetium Tc99m Pertechnate is excreted in human milk during lactation. It is not known whether Technetium Tc99m Sestamibi is excreted in human milk. Therefore, formula feedings should be substituted for breast feedings.

Pediatric Use

Safety and effectiveness in children below the age of 18 have not been established.

ADVERSE REACTIONS: During clinical trials, approximately 8% of patients experienced a transient metallic or bitter taste immediately after the injection of Technetium Tc99m Sestamibi. A few cases of transient headache, flushing and non-itching rash have also been attributed to administration of the agent. One patient demonstrated signs and symptoms consistent with seizure, 8 to 10 minutes after administration of the drug. No other adverse reactions specifically attributable to the use of Technetium Tc99m Sestamibi have been reported.

DOSAGE AND ADMINISTRATION: The suggested dose range for I.V. administration to be employed in the average patient (70 kg) is:

370 to 1110 MBq (10 to 30 mCi)

The dose administered should be the lowest required to provide an adequate study consistent with ALARA principles (See also PRECAUTIONS).

When used in the diagnosis of myocardial infarction, imaging should be completed within four hours after administration (see also CLINICAL PHARMACOLOGY section in full prescribing information).

The patient dose should be measured by a suitable radioactivity calibration system immediately prior to patient administration. Radiochemical purity should be checked prior to patient administration.

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration whenever solution and container permit.

Store at room temperature (15 to 30°) before and after reconstitution.

RADIATION DOSIMETRY: Table 4 shows the radiation doses to organs and tissues of an average patient (70 kg) per 1110 MBq (30 mCi) of Technetium Tc99m Sestamibi injected intravenously.

Table 4. Radiation Absorbed Doses from Tc99m Sestamibi

Organ	Estimated Radiation Absorbed Dose			
	REST			
	2.0 hour void		4.8 hour void	
	rads/ 30 mCi	mGy/ 1110 MBq	rads/ 30 mCi	mGy/ 1110 MBq
Breasts	0.2	2.0	0.2	1.9
Gallbladder Wall	2.0	20.0	2.0	20.0
Small Intestine	3.0	30.0	3.0	30.0
Upper Large Intestine Wall	5.4	55.5	5.4	55.5
Lower Large Intestine Wall	3.9	40.0	4.2	41.1
Stomach Wall	0.6	6.1	0.6	5.8
Heart Wall	0.5	5.1	0.5	4.9
Kidneys	2.0	20.0	2.0	20.0
Liver	0.6	5.8	0.6	5.7
Lungs	0.3	2.8	0.3	2.7
Bone Surfaces	0.7	6.8	0.7	6.4
Thyroid	0.7	7.0	0.7	6.8
Ovaries	1.5	15.5	1.6	15.5
Testes	0.3	3.4	0.4	3.9
Red Marrow	0.5	5.1	0.5	5.0
Urinary Bladder Wall	2.0	20.0	4.2	41.1
Total Body	0.5	4.8	0.5	4.8

Stabin, M., July, 1990, Oak Ridge Associated Universities, P.O. Box 117, Oak Ridge, TN 37831, (615) 576-3449.

HOW SUPPLIED: Du Pont's CARDIOLITE®, Kit for the preparation of Technetium Tc99m Sestamibi is supplied as a 5 mL vial in kits of two (2), five (5) and thirty (30) vials, sterile and non-pyrogenic.

Prior to lyophilization the pH is between 5.3 and 5.9. The contents of the vials are lyophilized and stored under nitrogen. Store at room temperature (15 to 30°C) before and after reconstitution. Technetium Tc99m Sestamibi contains no preservatives. Included in each two (2) vial kit is one (1) package insert, five (5) vial shield labels and five (5) radiation warning labels. Included in each five (5) vial kit is one (1) package insert, five (5) vial shield labels and five (5) radiation warning labels. Included in each thirty (30) vial kit is one (1) package insert, thirty (30) vial shield labels and thirty (30) radiation warning labels.

The US Nuclear Regulatory Commission has approved this reagent kit for distribution to persons licensed to use byproduct material identified in 35.100 and 35.200 of 10 CFR Part 35, to persons who hold an equivalent license issued by an Agreement State, and, outside the United States, to persons authorized by the appropriate authority.

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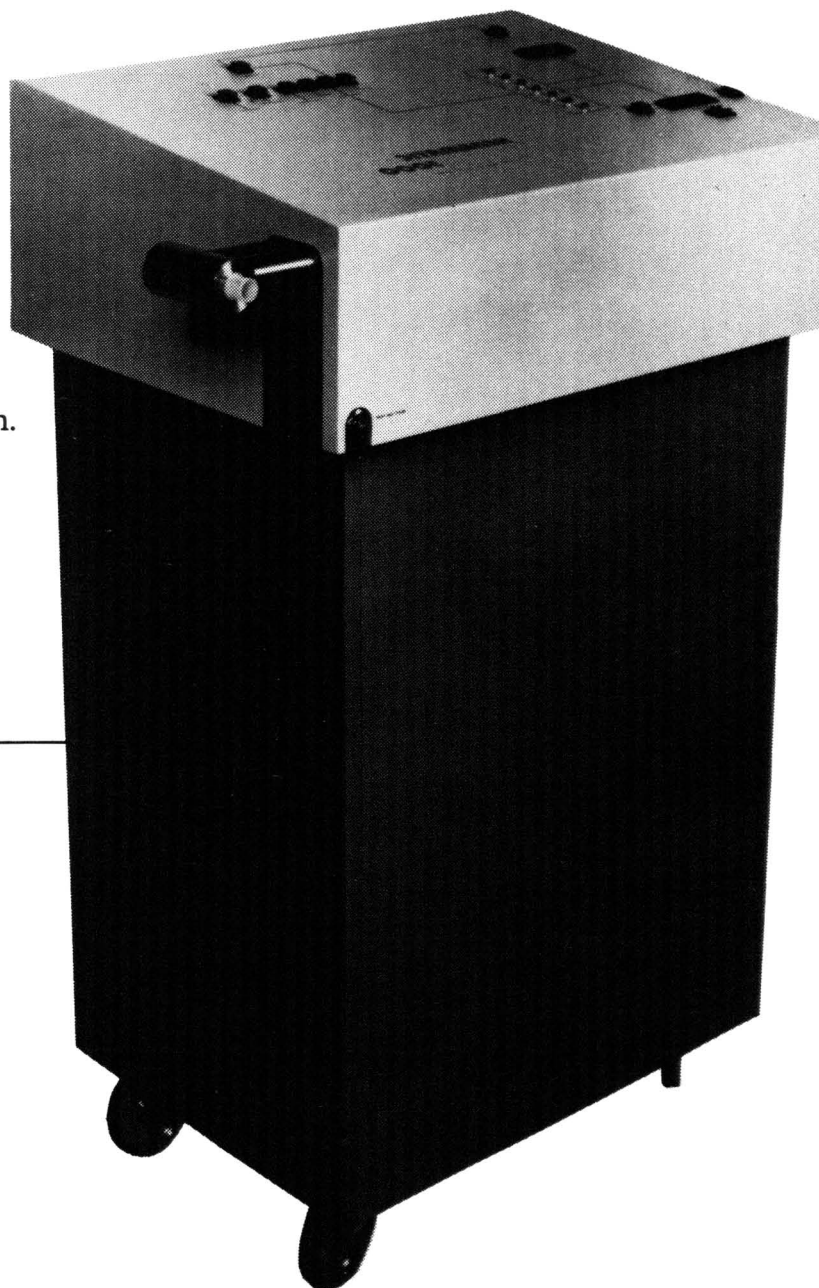
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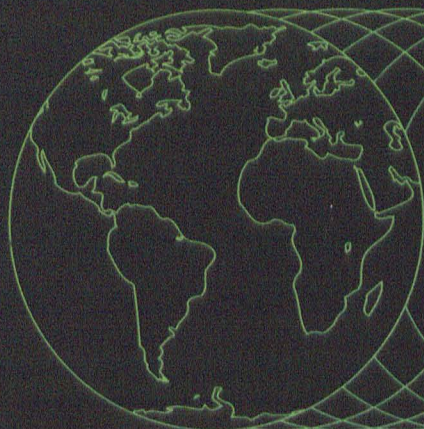
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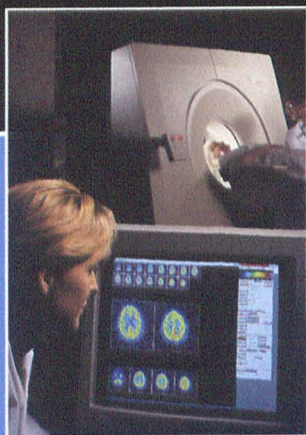


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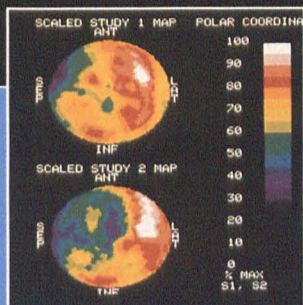


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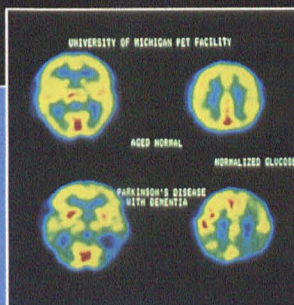
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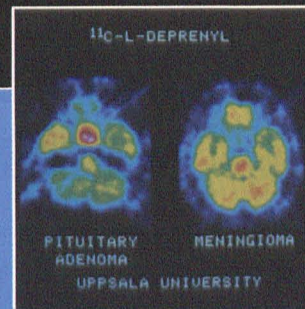
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CALL FOR ABSTRACTS

FOR Scientific Papers and Scientific Exhibits

1992 Scientific Program Committee, Scientific Exhibits Subcommittee, and the Scientific & Teaching Sessions Committee solicit the submission of abstracts from members and nonmembers of The Society of Nuclear Medicine for the 39th Annual Meeting in Los Angeles, CA. Scientific Paper abstracts accepted for the program will be published in a special supplement to the May issue of *The Journal of Nuclear Medicine* and accepted Technologist Section abstracts will be published in the June issue of the *Journal of Nuclear Medicine Technology*. Abstracts accepted for Society Program Scientific Exhibits will not be published. Original contributions on a variety of topics related to nuclear medicine will be considered, including:



The Society of
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Tuesday-Friday
June 9-12, 1992
Los Angeles, CA

Abstracts accepted for the program will be published in a special supplement to the May issue of *The Journal of Nuclear Medicine* and accepted Technologist Section abstracts will be published in the June issue of the *Journal of Nuclear Medicine Technology*. Abstracts accepted for Society Program Scientific Exhibits will not be published. Original contributions on a variety of topics related to nuclear medicine will be considered, including:

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- Instrumentation and Data Analysis
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- Immunology (antibody)
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- Hematology/Infectious Disease
- Oncology (non-antibody)

Authors seeking publication for the full text of their papers are strongly encouraged to submit their work for immediate review to the *JNM*, and for the technologist section, to the *JNMT*.

DEADLINES

For receipt of abstracts for
SCIENTIFIC PAPERS
is Tuesday, January 7, 1992.

For receipt of abstracts for
SCIENTIFIC EXHIBITS
is Tuesday, January 14, 1992.

There are two abstract forms for this year's meeting. The Scientific Paper abstract form can be obtained in the October 1991 *JNM*. The Scientific Exhibits abstract form is only available by calling or writing:

The Society of Nuclear Medicine
136 Madison Avenue,
Tel: (212) 889-0717

Att: Abstracts
New York, NY 10016-6760
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THE SOCIETY OF NUCLEAR MEDICINE MID-WINTER MEETING

Title: Computer and Instrumentation: Toward the 21st Century
Location: Hyatt Regency DFW, Dallas, TX
Date: Monday-Tuesday, February 10-11, 1992
Sponsor: The Computer and Instrumentation Council of The Society of Nuclear Medicine
CME Credit: Approximately 12 Hours AMA Category I
VOICE Credit: Approximately .9 CEUs available for VOICE Credit for Technologists
Seminar Notes: Registration includes a luncheon on Monday, February 10th, with a guest speaker. There are a limited amount of lunches available so please register early.

THE FEE	Before 12/20	On/After 12/20
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To make hotel reservations, call the Hyatt Regency DFW direct at (214) 453-1234. Indicate you are with The Society of Nuclear Medicine. Please make your reservations by January 10, 1992. Do NOT mail housing information to The Society.

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In November the
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If you have missed any of these forms,
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**SNM
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CRITICAL DATES

Item	Form Included in JNM	Due Date
ABSTRACT FORM Scientific Papers	October Issue	1/07/92
Scientific Exhibits	Contact SNM, Attn: Meetings Department	1/14/92
REGISTRATION FORM	November Issue	5/08/92
HOUSING FORM	December Issue	5/15/92

DON'T FORGET THE MID-WINTER MEETING IN DALLAS, TEXAS

DATE:

February 10-11, 1992

LOCATION:

Hyatt Regency DFW, Dallas, TX

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The Journal of Nuclear Medicine 1992 Reader Survey

You are invited to help shape the editorial policies of *The Journal of Nuclear Medicine*.

Please fill out the questionnaire and drop it in the mail. Our primary objective has always been to serve the needs of the nuclear medicine community—but unfortunately, the day-to-day operation of a biomedical journal often leaves little time for looking ahead, and no time for looking around. You can give that important perspective.

Give us a piece of your mind, and we'll give you a better Journal.

INSTRUCTIONS: The questions below may be answered by selecting letter codes from the selections beneath each question. Additional comments on separate sheets may be stapled to the form, but may require that additional postage be affixed. The *Journal* welcomes personal correspondence on any subject from readers and may be reached by phone at (617) 726-5785, or by fax at (617) 726-5708. Queries regarding subscription problems, government relations or matters not directly related to the basic editorial functions of the *Journal*, should be directed to the Society of Nuclear Medicine's offices in New York: (212) 889-0717.

1. How many hours **per month** do you spend reading the *Journal*?

- A. <1 B. 1-2 C. 2-3 D. >3

2. If the full text of *JNM* articles were available on-line, you would _____.

- A. Utilize the service at least once a week
B. Utilize the service at least once a month
C. Utilize less than once a month
D. Have no interest in this service

3. You are interested mainly in studies concerning: (choose 3, and number them in order of importance to you)

- A. Cardiology _____
B. Brain _____
C. Oncology _____
D. Respiratory _____
E. Hepatobiliary _____
F. Genitourinary _____
G. Endocrine _____
H. Gastroenterology _____
I. Skeletal _____
J. Antibodies _____
K. Instrumentation _____
L. Radiochemistry _____
M. Other _____

4. You would classify yourself as a(n) _____.

- A. Clinician
B. Technologist
C. Researcher
D. Industrial Representative
E. Regulator
F. Student

5. Which nuclear medicine procedures are performed most often in your department? (Again, please list your top three).

1. _____ 2. _____ 3. _____

6. You would like to see a greater number of _____ in the *Journal*.

- A. Case Reports
B. Basic Science Studies
C. Human Studies
D. Methodological Evaluations
E. Other _____

7. You would like to see a continuing medical education article on _____.

8. Your institution is located in _____.

- A. The United States or Canada
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F. Africa/Middle East

9. Rate the following *Journal* elements, from 1 to 10, for their value to you. 1 representing little or no value, 10 representing an element that you specifically look for each time you read the *Journal*.

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10. Considering your use of the *Journal*, your membership benefit of a subscription to *The Journal of Nuclear Medicine* is _____.

- A. a good value
B. in line with other journals
C. high, compared to other journals
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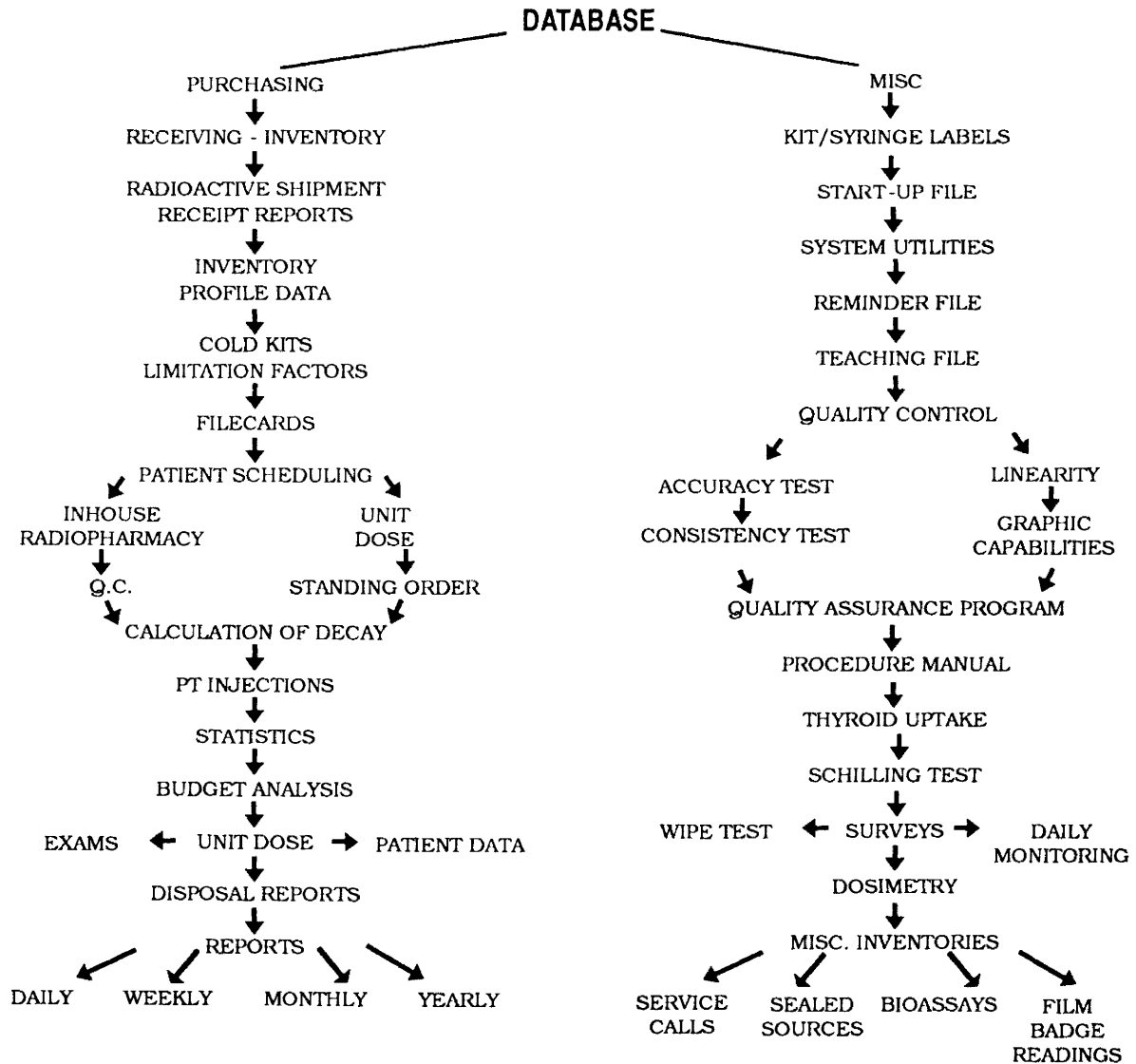
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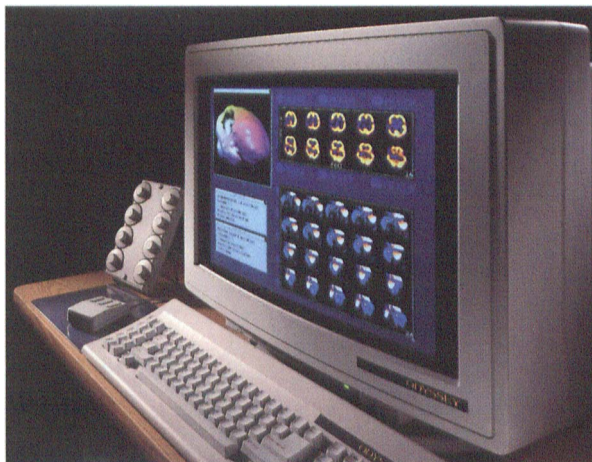
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PILOT RESEARCH AWARD

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I will need a _____ single/ _____ double room.

A check in the amount of \$650 should accompany this registration form and be made payable to the Medical College of Wisconsin. Telephone registrations must be confirmed by check within 10 days.

Name _____

Address _____

City/State/Zip _____

Office Phone (____) _____

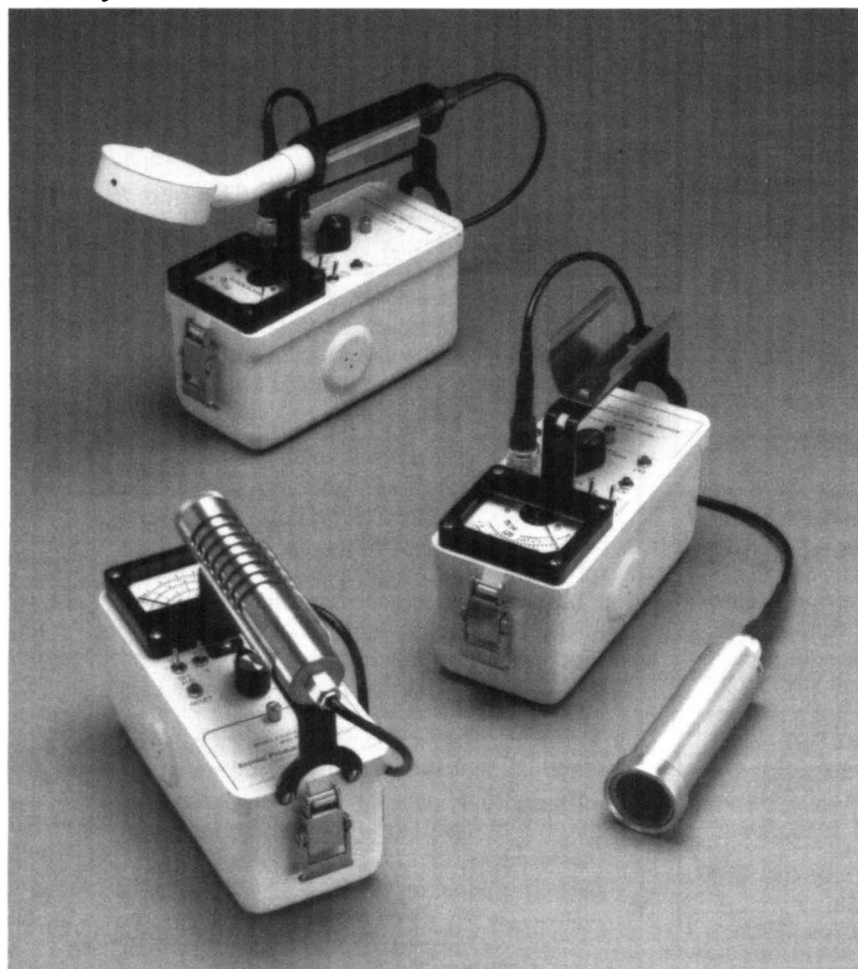
_____ work address _____ home address

Registrations and payment should be sent to:

LisaAnn Trembath
SPECT Brain Imaging Fellowship Coordinator
Nuclear Medicine Division
Medical College of Wisconsin
8700 W. Wisconsin Avenue
Milwaukee, WI 53226 (414) 257-7867

Each description of the products below was condensed from information supplied by the manufacturer. The reviews are published as a service to the professionals working in the field of nuclear medicine and their inclusion herein does not in any way imply an endorsement by the Editorial Board of The Journal of Nuclear Medicine or by The Society of Nuclear Medicine.

Survey Instruments



Atomic Products Corporation introduces a complete line of survey instruments including radiation meters, monitors, and probes. A key feature of the new RAM Survey Meter is its ability to interchange detectors without the need for recalibration. This saves valuable time while simultaneously ensuring accurate readings. The Model 2 Beta/Gamma Survey Meter comes complete with a built-in Side Wall GM Detector and telescoping metal

holder. Stray radiation is indicated by a built-in speaker and is ideal for measuring iodine-131, phosphorous-32, or higher energy beta radiation. The RAM Gamma Survey Meter is designed for one hand operation, utilizes only three pushbuttons, and provides fast time response and dead time correction. **Atomic Products Corporation, P.O. Box 702, Shirley, NY 11967. (516) 924-9000.**

Please see page 44A

High Resolution Graphics Board

XLI Corporation announces the availability of LaserPix 3.0 high-resolution graphics printer controller board. The unit, which

utilizes SUPER LGA Technology, offers true halftone print capability. Benchmark tests suggest that a test image can be produced in

less than 30 seconds using SUPER LGA technology. A full-page picture uses only half a megabyte of memory, compared with as much as 8 megabytes required by a dithered image. LaserPix upgrades standard 300 dpi laser printers to 2400 dpi (dots per inch equivalent) for photographic output. The product consists of two boards: the controller board is placed in the PC and the laser interface board resides in the printer. SUPER LGA technology allows the printer to vary the size of the individual dots that make up an image. Micrografx Picture Publisher is supplied free with LaserPix, providing image improvement tools such as grayscale adjustment, cropping, sizing, and scaling. Once the picture is edited, LaserPix is used to print the image. Images are produced with 256 gray levels at either 75- or 150-line screen, providing 2400 dpi. LaserPix is available immediately for all PC and compatible computers (Intel 80286 and higher) and industry standard laser printers with available video port. Print drivers are available for many popular Windows 3.0 applications including Aldus Pagemaker 4.0 and Ventura Publisher Windows edition and LaserPix supports PCL and Postscript printers. **XLI Corporation, 200 West Cummings Park, Suite 24, Woburn, MA 01801. (617) 932-9199 or (800) 338-0506.**

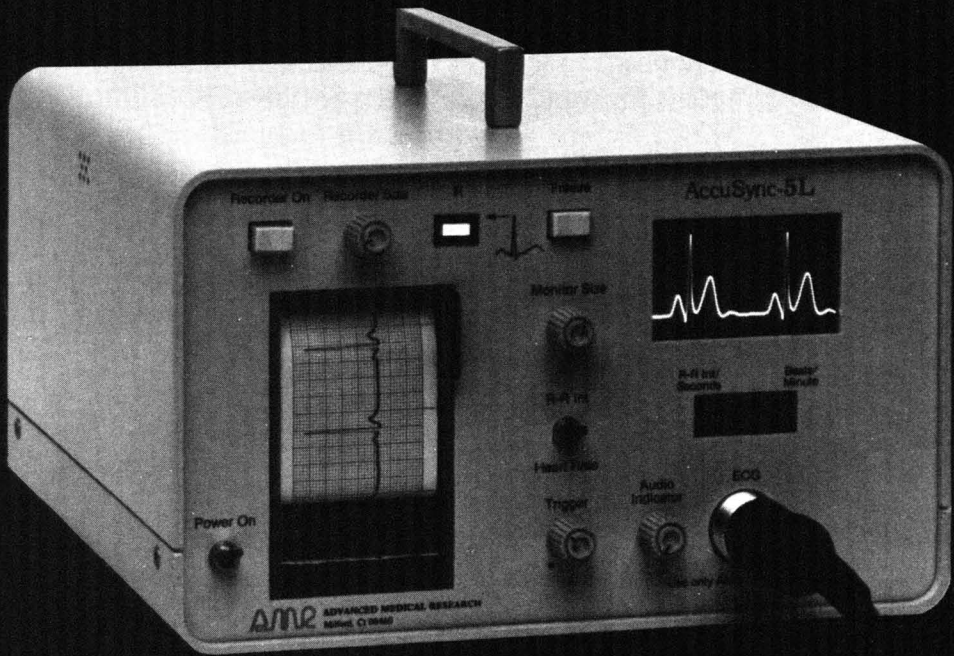
Please see page 44A

PLC Line Conditioner

Best Power Technology, Inc. introduces PLC Line Conditioners featuring surge suppression, true isolation, full voltage regulation, and harmonics suppression. PLC protects PCs, workstations, LAN nodes, and computer peripherals. For surge suppression capability, PLC can reduce a 250 volt surge to one volt. Both large and small surges never reach the load. PLC completely isolates its load from incoming utility power. An output neutral-to-ground bond eliminates the possibility of high-frequency line noise damaging the system. This design gives sensitive office equipment the best operating environment. PLC protects against many high and low-line voltage conditions. Only clean, regulated power reaches the load, even if the line is full of surges or sags. PLC also smooths out harmonic distortions. The unit's ferroresonant technology removes power line harmonics and gives a pure sine-wave output to the load to help protect equipment. A software PLC is available in sizes from 110 VA through 1.8 KVA. The hardware option fits needs between 2,000 and 15,000 watts. **Kenneth Urban, Best Power Technology, Inc., P.O. Box 280, Necedah, WI 54646. (608) 565-7200 or (800) 356-5794.**

Please see page 44A

AMR's AccuSync provides R-wave detection with precision and reliability.
The finest R-wave Triggering device available for computerized gated cardiac studies.

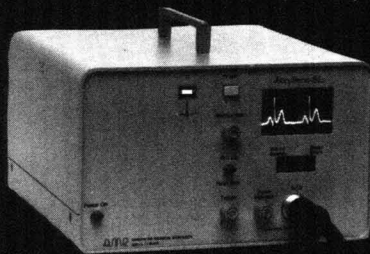


**AccuSync-5L
Features**

- Isolation Amplifier for Patient Safety
- Digital CRT Monitor
- ECG Strip Chart Recorder
- Heart Rate/R-R interval
- Trigger Pulse LED
- Trigger Control for Ease of Lead Placement and Precise Location of Trigger Pulse
- R-Trigger Output, Compatible with all Computers
- No Delay
- ECG Output
- Playback Mode (optional)
- Event Marker (optional)
- Audio Indicator

MODEL

AccuSync-6L



FEATURES

All **AccuSync-5L** features with the exception of the Strip Chart Recorder.

AccuSync-1L



All **AccuSync-5L** features with the exception of the Digital CRT Monitor.

AccuSync-3R



All **AccuSync-1L** features with the exception of the Strip Chart Recorder and Playback Mode.

AccuSync-4R



All **AccuSync-3R** features with the exception of the Heart Rate/R-R interval display.

JNM

DIRECT RESPONSE

The *Journal* is testing a new method to enable you to get information on a more timely basis from our advertisers.

Listed below are the companies that have advertised in this issue, as well as those that have been mentioned in the *New Products* section. Simply fill out the form and FAX it to the Society (FAX: 212/545-0221), and we will send it to the advertiser.

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Milford, CT
(203) 877-1610
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☐ Atomic Products
Shirley, NY
(516) 924-9000
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☐ Best Power Technology
Necedah, WI
(800) 356-5794
Page 44A

☐ Capintec, Inc.
Ramsey, NJ
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☐ Diversified Diagnostic
Products, Inc.
Houston, TX
(713) 955-5323
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☐ DuPont Company
No. Billerica, MA
(800) 343-7851
Following Page 8A

☐ Institute for Clinical PET
Washington, DC
(202) 466-4274
Pages 10A-11A

☐ Nuclear Associates
Carle Place, NY
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☐ Nuclear Medicine Consulting Firm
Greenville, PA
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☐ Siemens Medical Systems
Hoffman Estates, IL
(708) 304-7252
Pages IFC-1A

☐ Squibb Diagnostics
Princeton, NJ
(800) 257-5181
Pages IBC-OBC

☐ XLI Corporation
Woburn, MA
(800) 338-0506
Page 44A

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Circle One Answer In Each Category:

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 - a. 500 patients plus
 - b. 300-499 patients
 - c. 200-299 patients
 - d. 100-199 patients
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3. R&D Commercial
4. University
5. Government
6. Other _____

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 2. Specity
 3. Purchase
- #### Reason for Inquiry
1. Immediate Purchase
 2. General Information
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1. Yes
2. No

SNM Subscriber

1. Yes
2. No

CardioGen-82⁺
Rubidium Rb 82 Generator

INDICATIONS AND USAGE

Rubidium chloride Rb 82 injection is a myocardial perfusion agent that is useful in distinguishing normal from abnormal myocardium in patients with suspected myocardial infarction.

Cardiogen-82 (Rubidium Rb 82 Generator) must be used with an infusion system specifically labeled for use with the generator and capable of accurate measurement and delivery of doses of rubidium chloride Rb 82 injection not to exceed a single dose of 2220 MBq (60 mCi) and a cumulative dose of 4440 MBq (120 mCi) at a rate of 50 mL/min with a maximum volume per infusion of 100 mL and a cumulative volume not to exceed 200 mL. These performance characteristics reflect the conditions of use under which the drug development clinical trials were conducted.

Adequate data from clinical trials to determine precise localization of myocardial infarction or identification of stress-induced ischemia have not been collected.

Positron emission tomographic (PET) instrumentation is recommended for use with rubidium chloride Rb 82 injection.

CONTRAINDICATIONS

None known

WARNINGS

Caution should be used during infusion as patients with congestive heart failure may experience a transitory increase in circulatory volume load. These patients should be observed for several hours following the Rb-82 procedure to detect delayed hemodynamic disturbances.

PRECAUTIONS

General

Data are not available concerning the effect of marked alterations in blood glucose, insulin, or pH (such as is found in diabetes mellitus) on the quality of rubidium chloride Rb 82 scans. Attention is directed to the fact that rubidium is physiologically similar to potassium, and since the transport of potassium is affected by these factors, the possibility exists that rubidium may likewise be affected.

Rubidium chloride Rb 82 injection must be administered only with an appropriate infusion system capable of meeting the performance characteristics previously described (See **INDICATIONS AND USAGE**). The drug should be used only by those practitioners with a thorough understanding of the use and performance of the infusion system.

Repeat doses of rubidium chloride Rb 82 injection may lead to an accumulation of the longer lived radioactive contaminants strontium Sr 85 and strontium Sr 85.

Since eluate obtained from the generator is intended for intravenous administration, aseptic techniques must be strictly observed in all handling. Only additive free Sodium Chloride Injection USP should be used to elute the generator. Do not administer eluate from the generator if there is any evidence of foreign matter.

As in the use of any radioactive material, care should be taken to minimize radiation exposure to the patient consistent with proper patient management and to insure minimum radiation exposure to occupational workers.

Radiopharmaceuticals should be used only by physicians who are qualified by training and experience in the safe use and handling of radionuclides and whose experience and training have been approved by the appropriate government agency authorized to license the use of radionuclides.

Carcinogenesis, Mutagenesis, Impairment of Fertility

No long-term studies have been performed to evaluate carcinogenic potential, mutagenicity potential, or to determine whether rubidium Rb 82 may affect fertility in males or females.

Pregnancy Category C

Animal reproductive studies have not been conducted with rubidium Rb 82. It is also not known whether rubidium Rb 82 can cause fetal harm when administered to a pregnant woman or can affect reproductive capacity. Rubidium Rb 82 should be given to pregnant women only if the expected benefits to be gained clearly outweigh the potential hazards.

Ideally, examinations using radiopharmaceuticals, especially those examinations which are elective in nature, in women of childbearing capability should be performed during the first few (approximately 10) days following the onset of menses.

Nursing Mothers

It is not known whether rubidium Rb 82 is excreted in human milk. Due to the short half-life of rubidium Rb 82 (75 sec) it is unlikely that the drug would be excreted in human milk during lactation. However, because many drugs are excreted in human milk, caution should be exercised when rubidium Rb 82 is administered to nursing women.

Pediatric Use

Safety and effectiveness in children have not been established.

ADVERSE REACTIONS

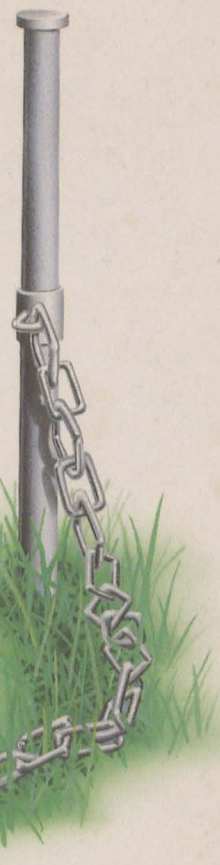
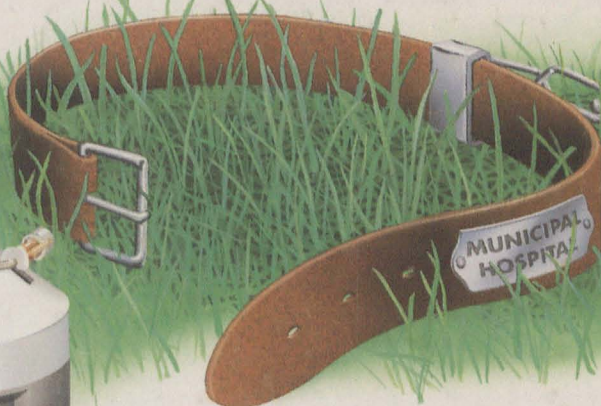
No adverse reactions specifically attributable to rubidium Rb 82 have been reported during controlled clinical trials.

HOW SUPPLIED

Cardiogen-82 (Rubidium Rb 82 Generator) is supplied in the form of strontium Sr 82 adsorbed on a hydrous stannic oxide column with an activity of 90-150 millicuries Sr-82 at calibration time. The generator is encased in a lead shield surrounded by a labeled plastic container. Complete assay data for each generator are provided on the container label. Cardiogen-82 (Rubidium Rb 82 Generator) is intended for use only with an appropriate, properly calibrated infusion system labeled for use with the generator.

(J4-263)

We've removed your PET collar



PET perfusion studies without a cyclotron

CardioGen-82® (Rubidium Rb 82 Generator) is the only generator-based myocardial perfusion agent indicated for PET imaging.

Now in 45 to 60 minutes you can have PET images to help you distinguish normal from abnormal myocardium. All without the expense of a cyclotron!

The short 75-second half-life lowers the radiation burden to the patient. When incorporated into the Rubidium Infusion System, serial imaging of myocardial blood flow changes can be performed as often as every ten minutes.



Rubidium-82
Infusion System

The CardioGen-82 System also improves patient throughput and scheduling efficiency by enabling you to perform multiple studies in a short time.

Remove the PET collar from your department. Get the PET images you need in 45 to 60 minutes, without a costly cyclotron.

CardioGen-82®
Rubidium Rb-82 Generator

Please see adjacent page for brief summary of prescribing information.

 **SQUIBB™**
Diagnostics